1=5

- 5. (Amended) An agent according to claim 2 in which the linker group is selected from an optionally substituted methylene chain, or -(CH<sub>2</sub>)<sub>m</sub>-Y-(CH<sub>2</sub>)<sub>n</sub>- wherein Y is selected from -O-, -S-, -S(O)-, -SO<sub>2</sub>-, -NH-, -Nalkyl-, -CO-, -OC(O)-, -NHC(O)-, -N(alkyl)C(O)-, -NHC(O)NH-, -NalkylC(O)NH-, NalkylC(O)Nalkyl-, -NHSO<sub>2</sub>-, -NalkylSO<sub>2</sub>-, NHSO<sub>2</sub>NH-, -NalkylSO<sub>2</sub>NH-, -NalkylSO<sub>3</sub>Nalkyl- and -OC(O)O-, m is 0-3 and n is 0-3.
- 6. (Amended) An agent according to claim 2 in which the nitric oxide synthase inhibitor moiety is selected from a group derived from an amino acid inhibitor of nitric oxide synthase a thiocitrulline derivative, an S-alkylisothiourea derivative or 2-aminopyridine derivative.

Please add the following new claims:

19. (New) An agent according to claim 3 in which the linker group X is a bond.

20. (New) An agent according to claim 3 in which the linker group is selected from an optionally substituted methylene chain, or -(CH<sub>2</sub>)<sub>m</sub>-Y-(CH<sub>2</sub>)<sub>n</sub>- wherein Y is selected from -O-, -S-, -S(O)-, -SO<sub>2</sub>-, -NH-, -Nalkyl-, -CO-, -OC(O)-, -NHC(O)-, -N(alkyl)C(O)-, -NHC(O)NH-, -NalkylC(O)NH-, NalkylC(O)Nalkyl-, -NHSO<sub>2</sub>-, -NalkylSO<sub>2</sub>-, -NHSO<sub>2</sub>NH-, -NalkylSO<sub>2</sub>NH-, -NalkylSO<sub>2</sub>Nalkyl- and -OC(O)O-, m is 0-3 and n is 0-3.